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1. 20080021226. 13 Oct 06. 24 Jan 08. Process for total synthesis of pladienolide B and pladienolide D. Kanada; Regina Mikie, et al. 549/215; 549/214 549/271 549/454 549/554 C07D303/02 20060101 C07D313/00 20060101 2. 20070155696. 28 Jan 05. 05 Jul 07. Method for stabilizing macrolide compounds. Ishihara; Hiroshi, et al. 514/58; 536/103 536/46 A61K31/724 20060101 C08B30/18 20060101 C08B3.7/16 20060101 3. 20060276339. 16 Oct 03. 07 Dec 06. Methods and compositions for increasing the efficacy of biologically-active ingredients. Windsor; J. Brian, et al. 504/127; 504/128 514/192 514/200 514/369 514/370 514/414 514/443 514/457 514/512 514/532 514/533 514/535 514/602 514/615 514/616 514/617 A01N43/16 20070101 A01N47/06 20070101 A01N57/00 20070101 A61K31/165 20070101 A61K31/18 20070101 A61K31/235 20070101 A61K31/24 20070101 A61K31/405 20070101 A61K31/425 20070101 A61K31/426 20070101 A61K31/43 20070101 A61K31/545 20070101 4. 20060241171. 23 Jun 06. 26 Oct 06. Novel physiologically active substances. Kotake; Yoshihiko, et al. 514/450; 549/271 A61K31/365 20060101 C07D313/04 20060101 C07D405/02 20060101 5. 20060235002. 31 Jul 03. 19 Oct 06. Novel physiologically active substance. Nagai; Mitsuo, et al. 514/217.03; 514/218 514/254.1 514/326 514/412 514/422 514/450 540/575 540/596 544/374 546/207 548/453 548/517 A61K31/4025 20060101 A61K31/407 20060101 A61K31/452 20060101 A61K31/496 20060101 A61K31/55 20060101 A61K31/551 20060101 C07D405/14 20060101 6. 20060141589. 27 Nov 03. 29 Jun 06. Method of producing macrolide compound. Okuda; Akifumi, et al. 435/123: 435/252.3 C12N1/21 20060101 C12P17/02 20060101 7. 20060079572. 30 Aug 05. 13 Apr 06. Novel bioactive substance. Mizui; Yoshiharu, et al. 514/450; 549/266 A61K31/365 20060101 C07D313/00 20060101 8. 20060009439. 29 May 03. 12 Jan 06. Novel physiologically active substances. Kotake; Yoshihiko, et al. 514/183; A61K31/33 20060101 9. 20050245514. 28 Jan 05. 03 Nov 05. Novel physiologically active substances. Kotake, Yoshihiko, et al. 514/232.5; 514/254.1 514/326 514/422 514/450 544/147 544/374 546/207 548/517 549/266 A61K031/5377 A61K031/496 A61K031/452 A61K031/4025 A61K031/365 C07D045/02 C07D413/02. 10. 20030077808. 17 Jan 01. 24 Apr 03. Nucleic acids, proteins, and antibodies. Rosen, Craig A., et al. 435/226; 435/320.1 435/325 435/69.1 435/69.4 530/399 536/23.1 C12N009/64 C07K014/575 C07H021/04 C12P021/02 C12N005/06. Print Generate Collection

Terms **Documents** L4 and 11107

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- 18. WO 200260890A. Physiologically-active Streptomyces-originated 12-membered ring macrolide compounds, useful in treating e.g. rheumatoid arthritis, angioma, inflammatory diseases, arthritis deformans, psoriasis. ASAI, N, et al. A61K031/335 A61K031/336 A61K031/365 A61K031/4025 A61K031/4427 A61K031/4523 A61K031/4545 A61K031/455 A61K031/496 A61K031/5375 A61K031/5377 A61K031/551 A61K031/74 A61P007/00 A61P009/00 A61P009/10 A61P017/06 A61P019/00 A61P019/02 A61P027/00 A61P027/02 A61P029/00 A61P035/00 A61P035/02 A61P035/04 A61P043/00 C07D313/00 C07D405/00 C07D405/14 C07D407/00 C07D407/06 C07D407/14 C07D493/00 C07D493/04 C07D493/08 C07D493/10 C12P017/02.

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L5: Entry 18 of 18

File: DWPI

Apr 19, 2007

DERWENT-ACC-NO: 2002-666923

DERWENT-WEEK: 200763

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TITLE: Physiologically-active Streptomyces-originated 12-membered ring macrolide compounds, useful in treating e.g. rheumatoid arthritis, angioma, inflammatory diseases, arthritis deformans, psoriasis.

#### Basic Abstract Text (1):

NOVELTY - Macrolide compounds (I), their pharmaceutically-acceptable salts or hydrates are new.

#### Basic Abstract Text (2):

DETAILED DESCRIPTION - Macrolide compounds of formula (I), their pharmaceuticallyacceptable salts or hydrates are new.

#### Basic Abstract Text (33):

(11) a process for producing the compounds, their pharmaceutically-accepta- ble salts or hydrates by using Streptomyces sp. (Mer-11107 FERM P-1844) or its mutant; and

#### Basic Abstract Text (45):

USE - The macrolide compounds are applicable in drug compositions to treat diseases requiring regulation of gene expression, inhibition of VEGF production or inhibition of neovascularization, or for treating solid tumors; or for treating angioma, inhibiting cancer metastasis, omental neovascularization, diabetic omentopathy, inflammatory diseases, arthritis deformans, rheumatoid arthritis, psoriasis, arteriosclerosis or solid tumors including cancer of the lung, brain tumor, breast cancer, prostate cancer, ovarian cancer, colon cancer and melanoma. (all claimed).

#### Standard Title Terms (1):

PHYSIOLOGICAL ACTIVE STREPTOMYCES ORIGIN MEMBER RING MACROLIDE COMPOUND USEFUL TREAT RHEUMATISM ARTHRITIS INFLAMMATION DISEASE ARTHRITIS PSORIASIS

First Hit

L5: Entry 2 of 18

File: PGPB

Jul 5, 2007

PGPUB-DOCUMENT-NUMBER: 20070155696

PGPUB-FILING-TYPE:

DOCUMENT-IDENTIFIER: US 20070155696 A1

TITLE: Method for stabilizing macrolide compounds

PUBLICATION-DATE: July 5, 2007

INVENTOR-INFORMATION:

CITY STATE COUNTRY NAME Ishihara; Hiroshi Ibaraki JP Takeda: Susumu Kumamoto JP JP Yamada; Tomonari Shizuoka Asahi; Yoshiaki Shizuoka JΡ

APPL-NO: 10/587042 [PALM] DATE FILED: January 28, 2005

FOREIGN-APPL-PRIORITY-DATA:

COUNTRY APPL-NO DOC-ID APPL-DATE

JP 2004-020804 2004JP-2004-020804 January 29, 2004

PCT-DATA:

DATE-FILED APPL-NO PUB-NO PUB-DATE 371-DATE

Jan 28, 2005 PCT/JP05/01637 Jul 24, 2006

INT-CL-PUBLISHED:

TYPE IPC DATE IPC-OLD
IPCP A61K31/724 20060101 A61K031/724
IPCS C08B3/18 20060101 C08B030/18
IPCS C08B37/16 20060101 C08B037/16

INT-CL-CURRENT:

TYPE IPC DATE
CIPP A61 K 31/724 20060101
CIPS C08 B 30/18 20060101
CIPS C08 B 37/16 20060101

US-CL-PUBLISHED: 514/058; 536/046, 536/103 US-CL-CURRENT: 514/58; 536/103, 536/46

ABSTRACT:

The present invention provides a method for stabilizing a macrolide compound, and an efficient method for producing the compound. Specifically, it provides a method for stabilizing a macrolide compound, in which a 12-membered ring macrolide compound, such as a compound expressed by the formula (1) and a cyclodextrin are both present, and a method for producing a macrolide compound, in which a cyclodextrin is made to be present in a culture broth of actinomycetes having an ability of producing the macrolide compound.

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New/Noteworthy 🚮 . E-Utilities	<u> </u>	Asai N, Kotake Y, Niijima J, Fukuda Y, Uehara T, Sakai T. Related Articles, Links
		Stereochemistry of pladienolide B.
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Clinical Queries Special Queries		Pladienolides, new substances from culture of Streptomyces platensis Mer- 11107. III. In vitro and in vivo antitumor activities.
LinkOut		J Antibiot (Tokyo). 2004 Mar;57(3):188-96.
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•		J Biol Chem. 2002 Mar 29;277(13):11107-15. Epub 2002 Jan 17. PMID: 11799110 [PubMed - indexed for MEDLINE]

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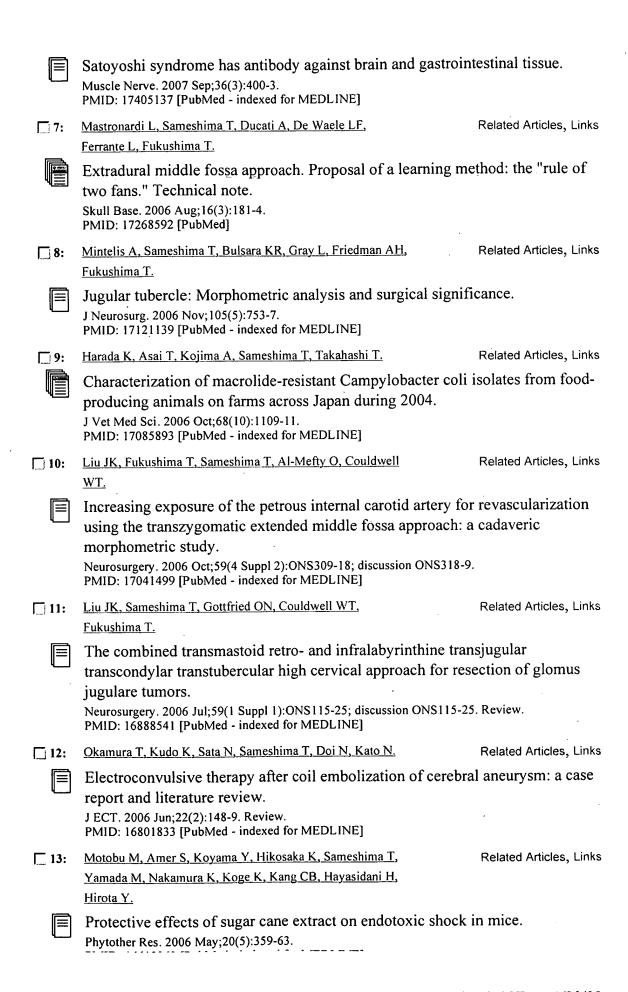
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